Increase of macrophage migration inhibitory factor in sera of patients with iridocyclitis

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Abstract

Aims—To determine whether macrophage migration inhibitory factor (MIF) levels were increased in sera of the patients with iridocyclitis.

Methods—Sera were obtained from 41 patients with acute iridocyclitis, 13 patients with chronic iridocyclitis, and 44 healthy control subjects. MIF levels were determined by a human MIF ELISA.

Results—The average levels of MIF in the sera of patients with both acute and chronic iridocyclitis were significantly higher than that of healthy subjects.

Conclusion—Uveitis induces the elevation of serum MIF, which may affect various inflammatory symptoms in uveitis.

Patients and methods

After informed consent was obtained, sera were obtained from 41 age and sex matched patients with acute iridocyclitis (13 patients have HLA-B27), 13 patients with chronic iridocyclitis, and 44 healthy control subjects. Acute cases of iridocyclitis had a sudden onset which lasted up to 8 weeks. Chronic forms of iridocyclitis had an insidious onset and lasted longer than 8 weeks. Systemic symptoms as seen in ankylosing spondylitis (AS) and other collagen diseases were not complicated in these patients. Infectious iridocyclitis, Fuch's heterochromia, and Posner–Schlossman syndrome were not included in this study. Blood samples were collected at the first medical examination in the uveitis survey clinic of Hokkaido University Hospital. All patients were in active phase of iridocyclitis when blood samples were collected. All of the patients with acute and chronic iridocyclitis were given topical corticosteroids at this time. MIF levels were determined by a human MIF enzyme linked immunosorbent assay (ELISA) (Cosmobio, Tokyo, Japan) as described previously. Statistical analysis was performed using the Mann–Whitney U test. This study was performed according to the tenets of the Declaration of Helsinki.

Results

The mean MIF concentrations in the sera of patients with acute iridocyclitis, idiopathic chronic iridocyclitis, and healthy control subjects were 28.8 (SD 3.4), 18.5 (5.9), and 4.4 (0.2) ng/ml, respectively. The average level of MIF in the sera of iridocyclitis patients was significantly higher (p<0.0001) than that of healthy control subjects (Fig 1). No significant difference was found in the serum MIF level between the patients with acute iridocyclitis and those of chronic iridocyclitis. However, since only 13 samples were evaluated from the chronic group in the present study, the two high MIF samples might significantly alter the distribution. Examination of more samples from cases of chronic iridocyclitis is needed to confirm this result.

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In the present study, we detected considerably high levels of serum MIF in the patients with iridocyclitis who showed no systemic inflammation. This finding suggests that possible origins of MIF in patients with uveitis. The first source may be the uvea of these patients, including those with Behçet’s disease and sarcoidosis. In previous studies, we observed that high levels of MIF were present in uvea\(^1\) and vascular endothelial cells.\(^2\) Thus, it seems highly likely that a various cells in the ocular are responsible for MIF production. The second source may be the pituitary gland that is stimulated by ocular inflammation and releases MIF in the serum. In the murine model, it was reported that MIF was circulated normally as a hormone and was released in part by the pituitary gland in response to stress or systemic inflammatory stimuli.\(^3\) \(^4\) The third source may be peripheral blood mononuclear cells (PBMC). Recently, Shimizu \textit{et al} reported that MIF production by PBMC was markedly upregulated in patients with atopic dermatitis.\(^5\) It seems that MIF produced in ocular tissues of iridocyclitis are in some way related to the systemic production of MIF via the pituitary gland, PBMC, and other cells even though apparent systemic inflammatory symptoms are not observed in patients with iridocyclitis. We would like to emphasise here the close association between uveitis and MIF in humans.

We and Bacher \textit{et al} reported that anti-MIF Ab inhibited \textit{T} cell proliferative responses in vitro.\(^6\) \(^7\) Furthermore, we recently reported that anti-MIF mAb administration inhibited experimental autoimmune uveoretinitis (EAU) in the murine model.\(^8\) Thus, MIF produced by various cells appears to be involved in the enhancement of ocular inflammation and immune responses.

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